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## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification 7 :</b> <b>C07K 14/435, C12N 1/21, 5/10, 15/12, 15/67, 15/70, 15/85</b>	<b>A1</b>	<b>(11) International Publication Number:</b> <b>WO 00/61620</b> <b>(43) International Publication Date:</b> 19 October 2000 (19.10.00)
<b>(21) International Application Number:</b> PCT/US00/09069 <b>(22) International Filing Date:</b> 6 April 2000 (06.04.00) <b>(30) Priority Data:</b> 60/128,702 9 April 1999 (09.04.99) US 60/177,049 20 January 2000 (20.01.00) US <b>(71) Applicant (for all designated States except US):</b> HUMAN GENOME SCIENCES, INC. [US/US]; 9410 Key West Avenue, Rockville, MD 20850 (US). <b>(71)(72) Applicant and Inventor:</b> ROSEN, Craig, A. [US/US]; 22400 Rolling Hill Road, Laytonsville, MD 20882 (US). <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only):</b> RUBEN, Steven, M. [US/US]; 18528 Heritage Hills Drive, Laytonsville, MD 20882 (US). KOMATSOUKIS, George [US/US]; 9518 Garwood Street, Silver Spring, MD 20901 (US). <b>(74) Agents:</b> HOOVER, Kenley, K. et al.; Human Genome Sciences, Inc., 9410 Key West Avenue, Rockville, MD 20850 (US).		<b>(81) Designated States:</b> AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).  <b>Published</b> <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
<b>(54) Title:</b> 49 HUMAN SECRETED PROTEINS  <b>(57) Abstract</b>  The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.		

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EE	Estonia						

nucleotide residues shown in SEQ ID NO:43, and where b is greater than or equal to a + 14.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 34

- 5           The computer algorithm BLASTX has been used to determine that the translation product of this gene shares sequence homology with, as a non-limiting example, the sequence accessible through the following database accession no. gi|57671 (all information available through the recited accession number is incorporated herein by reference) which is described therein as "ribonuclease inhibitor [Rattus norvegicus]." A partial alignment demonstrating the observed
- 10           homology is shown immediately below.

```
>gi|57671 ribonuclease inhibitor [Rattus norvegicus] >pir|S20597|S20597
      ribonuclease inhibitor - rat
      Length = 456

      Plus Strand HSPs:

      Score = 519 (182.7 bits), Expect = 3.6e-49, P = 3.6e-49
      Identities = 123/325 (37%), Positives = 170/325 (52%), Frame = +1

      Q: 640 EMGLAINDSFLSASLXRILCEQIASDTCHLQRVVFKNISPADAHRNLCL-ALRGHKTVTY 816
          E+ L N+ A + +L + + + TC +Q++ +N S +A + LR T+
      S: 55 ELSLRTNE-LGDAGVGLVL-QGLQNPTCKIQKLSLQNCSLTEAGCGVLPDVLRSLSLIRE 112

      Q: 817 LTLOGNDQ-DDMFPALCEVLRHPECNLRYLGLVSCSATTQOWADLSLALEVNQSLTCVNL 993
          L L N D+ LCE LR P+C L L L C+ T L+ L V + L
      S: 113 LHLNDNPLGDEGLKLLCEGLRDPQCRLEKQLLEYCNLTATSCEPLASVLRVKPDFKELVL 172

      Q: 994 SDNELLDEGAKLLYTTLRHPKCFLOQLSLENCHLTEANCKDLAAVLVVSRELTHLCLAKN 1173
          S+N+ + G L L+ C L+ L LENC +T ANCKDL V+ L L L N
      S: 173 SNNDPFHEAGIHTLCQGLKDSACQLESRLKLENCGITSANCKDLCDVVASKASLQELDLSN 232

      Q: 1174 FIGNTGVKFLCEGLRYPECKLQTLVLWNC DITS DGCCDLTKLLQEKSSLLCLDLGLNHIG 1353
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      Q: 1354 VKGMKFLCEALRKPLCNLRCLWLWGCSIPFSCEDLCSALSCNQSLVTLDLGQNPLGSSG 1533
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      S: 293 DEGAQLLCESLLEPGCQLESILVVKTCSLTAASCPHFCSVLTKNSSLFELQMSSNPLGDSG 352

      Q: 1534 VKMLFETLTCSSGTLRTLRLKIDDFND 1614
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V L + L      L R L L    D    D  
 S:    353 VVELCKALGYPDVLRVLWLGDGCDVTD 379

The segment of gil57671 that is shown as "S" above is set out in the sequence listing as SEQ ID NO. 143. Based on the structural similarity these homologous polypeptides are expected to share at least some biological activities. Such activities are known in the art, some of which are described elsewhere herein. Assays for determining such activities are also known in the art, some of which have been described elsewhere herein.

Preferred polypeptides of the invention comprise a polypeptide having the amino acid sequence set out in the sequence listing as SEQ ID NO. 144 which corresponds to the "Q" sequence in the alignment shown above (gaps introduced in a sequence by the computer are, of course, removed).

It has been discovered that this gene is expressed primarily in the following tissues/cDNA libraries: Human Testes Tumor and to a lesser extent in Soares fetal liver spleen 1NFLS; NCI\_CGAP\_GCB1; Human Testes; Human Endometrial Tumor; Soares placenta Nb2HP; Soares infant brain 1NIB; 12 Week Old Early Stage Human, II; Human Uterine Cancer; Human Whole Six Week Old Embryo; Activated T-Cell (12hs)/Thiouridine labelledEco; Spleen, Chronic lymphocytic leukemia and Soares\_placenta\_8to9weeks\_2NbHP8to9W.

Preferred epitopes include those comprising a sequence shown in SEQ ID NO. 93 as residues: Asp-194 to Leu-199, Ile-206 to Pro-211, Glu-224 to Ser-229.

Many polynucleotide sequences, such as EST sequences, are publicly available and accessible through sequence databases. Some of these sequences are related to SEQ ID NO:44 and may have been publicly available prior to conception of the present invention. Preferably, such related polynucleotides are specifically excluded from the scope of the present invention. To list every related sequence

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would be cumbersome. Accordingly, preferably excluded from the present invention are one or more polynucleotides comprising a nucleotide sequence described by the general formula of a-b, where a is any integer between 1 to 1978 of SEQ ID NO:44, b is an integer of 15 to 1992, where both a and b correspond to the positions of  
 5 nucleotide residues shown in SEQ ID NO:44, and where b is greater than or equal to a + 14.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 35

The computer algorithm BLASTX has been used to determine that the  
 10 translation product of this gene shares sequence homology with, as a non-limiting example, the sequence accessible through the following database accession no. gi|2559012 (all information available through the recited accession number is incorporated herein by reference) which is described therein as "chaperonin containing t-complex polypeptide 1, beta subunit; CCT-beta [Homo sapiens]." A  
 15 partial alignment demonstrating the observed homology is shown immediately below.

```

    >gi|2559012 (AF026293) chaperonin containing t-complex polypeptide 1, beta
    subunit; CCT-beta [Homo sapiens] >gi|2559012 (AF026293)
  20 chaperonin
    containing t-complex polypeptide 1, beta subunit; CCT-beta
    [Homo
    sapiens] >gi|4090929 (AF026166) chaperonin-containing TCP-1
    beta
    subunit homolog [Homo sapiens] >sp|G4090929|G4090929
  25 CHAPERONIN-CONTAINING TCP-1 BETA SUBUNIT HOMOLOG.
    >sp|G2559012|G2559012 CHAPERONIN CONTAINING T-COMPLEX
    POLYPEPTIDE
    1, BETA SUBUNIT. >gi|1871210 T-complex protein 1, Beta subunit
    (TCP-1-BETA) [Homo sapiens] {SUB 1-217}
  30 Length = 535
  
```

#### Plus Strand HSPs:

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  35 Score = 2610 (918.8 bits), Expect = 9.4e-271, P = 9.4e-271
    Identities = 525/535 (98%), Positives = 525/535 (98%), Frame = +2
  
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Q: 92 MASLSLAPVNIFKAGADEERAETARLTSTFIGAIAIGDLVKSTLGPKGMDRILLSSGRDAS 271

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***What Is Claimed Is:***

1. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of:
  - (a) a polynucleotide fragment of SEQ ID NO:X or a polynucleotide fragment of the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
  - (b) a polynucleotide encoding a polypeptide fragment of SEQ ID NO:Y or a polypeptide fragment encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
  - (c) a polynucleotide encoding a polypeptide domain of SEQ ID NO:Y or a polypeptide domain encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
  - 15 (d) a polynucleotide encoding a polypeptide epitope of SEQ ID NO:Y or a polypeptide epitope encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
  - (e) a polynucleotide encoding a polypeptide of SEQ ID NO:Y or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X, 20 having biological activity;
  - (f) a polynucleotide which is a variant of SEQ ID NO:X;
  - (g) a polynucleotide which is an allelic variant of SEQ ID NO:X;
  - (h) a polynucleotide which encodes a species homologue of the SEQ ID NO:Y;

(i) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.

5

2. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding a secreted protein.

10

3. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding the sequence identified as SEQ ID NO:Y or the polypeptide encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.

15

4. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises the entire nucleotide sequence of SEQ ID NO:X or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.

20

5. The isolated nucleic acid molecule of claim 2, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.

25

6. The isolated nucleic acid molecule of claim 3, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.

7. A recombinant vector comprising the isolated nucleic acid molecule of claim 1.

5 8. A method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 1.

9. A recombinant host cell produced by the method of claim 8.

10 10. The recombinant host cell of claim 9 comprising vector sequences.

11. An isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from the group consisting of:

(a) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence  
15 included in ATCC Deposit No:Z;

(b) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z, having biological activity;

(c) a polypeptide domain of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

20 (d) a polypeptide epitope of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

(e) a secreted form of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

(f) a full length protein of SEQ ID NO:Y or the encoded sequence included in  
25 ATCC Deposit No:Z;

(g) a variant of SEQ ID NO:Y;

(h) an allelic variant of SEQ ID NO:Y; or

(i) a species homologue of the SEQ ID NO:Y.

12. The isolated polypeptide of claim 11, wherein the secreted form or the full length protein comprises sequential amino acid deletions from either the C-terminus or the N-terminus.

13. An isolated antibody that binds specifically to the isolated polypeptide of claim 11.

10 14. A recombinant host cell that expresses the isolated polypeptide of claim 11.

15. A method of making an isolated polypeptide comprising:

(a) culturing the recombinant host cell of claim 14 under conditions such that said polypeptide is expressed; and  
15 (b) recovering said polypeptide.

16. The polypeptide produced by claim 15.

20 17. A method for preventing, treating, or ameliorating a medical condition, comprising administering to a mammalian subject a therapeutically effective amount of the polypeptide of claim 11 or the polynucleotide of claim 1.

25 18. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:

(a) determining the presence or absence of a mutation in the polynucleotide of claim 1; and

(b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or absence of said mutation.

5

19. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:

(a) determining the presence or amount of expression of the polypeptide of claim 11 in a biological sample; and

10 (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or amount of expression of the polypeptide.

20. A method for identifying a binding partner to the polypeptide of claim 11 comprising:

15 (a) contacting the polypeptide of claim 11 with a binding partner; and  
(b) determining whether the binding partner effects an activity of the polypeptide.

21. The gene corresponding to the cDNA sequence of SEQ ID NO:Y.

20

22. A method of identifying an activity in a biological assay, wherein the method comprises:

(a) expressing SEQ ID NO:X in a cell;

(b) isolating the supernatant;

25 (c) detecting an activity in a biological assay; and

(d) identifying the protein in the supernatant having the activity.

23. The product produced by the method of claim 20.

&lt;400&gt; 43

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&lt;210&gt; 44

&lt;211&gt; 1992

&lt;212&gt; DNA

&lt;213&gt; Homo sapiens

&lt;400&gt; 44

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1992

&lt;210&gt; 45

&lt;211&gt; 1973

&lt;212&gt; DNA

&lt;213&gt; Homo sapiens

&lt;400&gt; 45

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&lt;210&gt; 46

&lt;211&gt; 1924

&lt;212&gt; DNA

&lt;213&gt; Homo sapiens

&lt;400&gt; 46

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50

55

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<220>  
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&lt;400&gt; 92

Met Ile Thr Asp Val Gln Leu Ala Ile Phe Ala Asn Met Leu Gly Val  
 1 5 10 15

Ser Leu Phe Leu Leu Val Val Leu Tyr His Tyr Val Ala Val Asn Asn  
 20 25 30

Pro Lys Lys Gln Glu Xaa  
 35

<210> 93  
 <211> 235  
 <212> PRT  
 <213> Homo sapiens

<220>  
 <221> SITE  
 <222> (235)  
 <223> Xaa equals stop translation

&lt;400&gt; 93

Lys Leu Leu Tyr Thr Thr Leu Arg His Pro Lys Cys Phe Leu Gln Arg  
 1 5 10 15

Leu Ser Leu Glu Asn Cys His Leu Thr Glu Ala Asn Cys Lys Asp Leu  
 20 25 30

Ala Ala Val Leu Val Val Ser Arg Glu Leu Thr His Leu Cys Leu Ala  
 35 40 45

Lys Asn Pro Ile Gly Asn Thr Gly Val Lys Phe Leu Cys Glu Gly Leu  
 50 55 60

Arg Tyr Pro Glu Cys Lys Leu Gln Thr Leu Val Leu Trp Asn Cys Asp  
 65 70 75 80

Ile Thr Ser Asp Gly Cys Cys Asp Leu Thr Lys Leu Leu Gln Glu Lys

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66

85								90						95					
Ser	Ser	Leu	Leu	Cys	Leu	Asp	Leu	Gly	Leu	Asn	His	Ile	Gly	Val	Lys				
				100								105							
Gly	Met	Lys	Phe	Leu	Cys	Glu	Ala	Leu	Arg	Lys	Pro	Leu	Cys	Asn	Leu				
				115								120							
Arg	Cys	Leu	Trp	Leu	Trp	Gly	Cys	Ser	Ile	Pro	Pro	Phe	Ser	Cys	Glu				
				130								135							
Asp	Leu	Cys	Ser	Ala	Leu	Ser	Cys	Asn	Gln	Ser	Leu	Val	Thr	Leu	Asp				
				145								150							
Leu	Gly	Gln	Asn	Pro	Leu	Gly	Ser	Ser	Gly	Val	Lys	Met	Leu	Phe	Glu				
				165								170							
Thr	Leu	Thr	Cys	Ser	Ser	Gly	Thr	Leu	Arg	Thr	Leu	Arg	Leu	Lys	Ile				
				180								185							
Asp	Asp	Phe	Asn	Asp	Glu	Leu	Asn	Lys	Leu	Leu	Glu	Glu	Ile	Glu	Glu				
				195								200							
Lys	Asn	Pro	Gln	Leu	Ile	Ile	Asp	Thr	Glu	Lys	His	His	Pro	Trp	Glu				
				210								215							
Glu	Arg	Pro	Ser	Ser	His	Asp	Phe	Met	Ile	Xaa									
				225								230							
												235							

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<210> 94
<211> 9
<212> PRT
<213> Homo sapiens
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<220>
<221> SITE
<222> (9)
<223> Xaa equals stop translation
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<400> 94  
Ser Leu Phe Trp Gly Cys Pro Trp Xaa  
1 5

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<210> 95
<211> 42
<212> PRT
<213> Homo sapiens
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**<220>**

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Glu Ser Leu Trp Val Lys Thr Cys Ser Leu Thr Ala Ala Ser Cys Pro  
 260 265 270

His Phe Cys Ser Val Leu Thr Lys Asn Ser Ser Leu Phe Glu Leu Gln  
 275 280 285

Met Ser Ser Asn Pro Leu Gly Asp Ser Gly Val Val Glu Leu Cys Lys  
 290 295 300

Ala Leu Gly Tyr Pro Asp Thr Val Leu Arg Val Leu Trp Leu Gly Asp  
 305 310 315 320

Cys Asp Val Thr Asp  
 325

<210> 144

<211> 325

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (16)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 144

Glu Met Gly Leu Ala Ile Asn Asp Ser Phe Leu Ser Ala Ser Leu Xaa  
 1 5 10 15

Arg Ile Leu Cys Glu Gln Ile Ala Ser Asp Thr Cys His Leu Gln Arg  
 20 25 30

Val Val Phe Lys Asn Ile Ser Pro Ala Asp Ala His Arg Asn Leu Cys  
 35 40 45

Leu Ala Leu Arg Gly His Lys Thr Val Thr Tyr Leu Thr Leu Gln Gly  
 50 55 60

Asn Asp Gln Asp Asp Met Phe Pro Ala Leu Cys Glu Val Leu Arg His  
 65 70 75 80

Pro Glu Cys Asn Leu Arg Tyr Leu Gly Leu Val Ser Cys Ser Ala Thr  
 85 90 95

Thr Gln Gln Trp Ala Asp Leu Ser Leu Ala Leu Glu Val Asn Gln Ser  
 100 105 110

Leu Thr Cys Val Asn Leu Ser Asp Asn Glu Leu Leu Asp Glu Gly Ala  
 115 120 125

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Lys Leu Leu Tyr Thr Thr Leu Arg His Pro Lys Cys Phe Leu Gln Arg  
 130 135 140  
 Leu Ser Leu Glu Asn Cys His Leu Thr Glu Ala Asn Cys Lys Asp Leu  
 145 150 155 160  
 Ala Ala Val Leu Val Val Ser Arg Glu Leu Thr His Leu Cys Leu Ala  
 165 170 175  
 Lys Asn Pro Ile Gly Asn Thr Gly Val Lys Phe Leu Cys Glu Gly Leu  
 180 185 190  
 Arg Tyr Pro Glu Cys Lys Leu Gln Thr Leu Val Leu Trp Asn Cys Asp  
 195 200 205  
 Ile Thr Ser Asp Gly Cys Cys Asp Leu Thr Lys Leu Leu Gln Glu Lys  
 210 215 220  
 Ser Ser Leu Leu Cys Leu Asp Leu Gly Leu Asn His Ile Gly Val Lys  
 225 230 235 240  
 Gly Met Lys Phe Leu Cys Glu Ala Leu Arg Lys Pro Leu Cys Asn Leu  
 245 250 255  
 Arg Cys Leu Trp Leu Trp Gly Cys Ser Ile Pro Pro Phe Ser Cys Glu  
 260 265 270  
 Asp Leu Cys Ser Ala Leu Ser Cys Asn Gln Ser Leu Val Thr Leu Asp  
 275 280 285  
 Leu Gly Gln Asn Pro Leu Gly Ser Ser Gly Val Lys Met Leu Phe Glu  
 290 295 300  
 Thr Leu Thr Cys Ser Ser Gly Thr Leu Arg Thr Leu Arg Leu Lys Ile  
 305 310 315 320  
 Asp Asp Phe Asn Asp  
 325

&lt;210&gt; 145

&lt;211&gt; 535

&lt;212&gt; PRT

&lt;213&gt; Homo sapiens

&lt;400&gt; 145

Met Ala Ser Leu Ser Leu Ala Pro Val Asn Ile Phe Lys Ala Gly Ala  
 1 5 10 15

Asp Glu Glu Arg Ala Glu Thr Ala Arg Leu Thr Ser Phe Ile Gly Ala

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